Open Peer Review on Qeios

Roberts syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Roberts syndrome. ORPHA:3103

Roberts syndrome (RBS) is characterized by pre- and postnatal growth retardation, severe symmetric limb reduction defects, craniofacial anomalies and severe intellectual deficit. SC phocomelia is a milder form of RBS.